Clean copy of pending claims as amended in this response:

1. A stable liquid pharmaceutical botulinum toxin formulation, comprising a pharmaceutically acceptable buffer capable of providing a buffered pH range between about pH 5 and pH 6, and

isolated botulinum toxin;

wherein said formulation is stable as a liquid for at least one year at a temperature between about 0 and 10 degrees centigrade.

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- 2. The formulation of claim 1, wherein said temperature is about 5±3 degrees centigrade.
- 3. The formulation of claim 1, wherein said temperature is about 4±2 degrees centigrade.
 - 4. The formulation of claim 1, wherein said buffered pH range is about pH 5.6±0.2.
- 5. The formulation of claim 1, wherein said toxin formulation is stable in liquid form for at least two years.
- 6. The formulation of claim 1, wherein said buffer has a pK in the range of pH 4.5-6.5.
- 7. The formulation of claim 6, wherein said buffer is selected from the group consisting of phosphate buffer, phosphate-citrate buffer, and succinate buffer.

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8. The formulation of claim 1, wherein said botulinum toxin is of a botulinum toxin type selected from the group consisting of Types A, B, C_1 , C_2 , D, E, F and G.

- 9. The formulation of claim 8, wherein said botulinum toxin is botulinum toxin Type B present at a concentration in the range of about 100-20,000 U/ml.
- 10. The formulation of claim 9, wherein said botulinum toxin Type B is present in a high molecular weight complex of about 700 kilodaltons (kD).
- 11. The formulation of claim 9, wherein said botulinum toxin Type B is present at a concentration between about 1000-5000 U/ml.
- 12. The formulation of claim 8, wherein said botulinum toxin is botulinum toxin Type A, present at a concentration in the range of about 20-2000 U/ml.
- 13. The formulation of claim 12, wherein said botulinum toxin Type A is present at a concentration in the range of about 100-1000 U/ml.
 - 14. The formulation of claim 1, which further includes an excipient protein.
- 15. The formulation of claim 14, wherein said excipient protein is selected from the group consisting of serum albumin, recombinant human serum albumin, and gelatin.
- 16. A stable liquid pharmaceutical botulinum toxin formulation, comprising a pharmaceutically acceptable liquid buffer capable of providing a buffered pH range between about pH 5 and pH 6, and

isolated botulinum toxin;

wherein said toxin formulation is stable as a liquid for at least about 6 months at a temperature between about 10 and 30 degrees centigrade.

- 17. The formulation of claim 16, wherein said temperature is about 25°C.
- 18. The formulation of claim 16, wherein said buffered pH range is about pH 5.6±0.2.

- 19. The formulation of claim 16, wherein said buffer has a pK in the range of pH 4.5-6.5.
- 20. The formulation of claim 19, wherein said buffer is selected from the group consisting of phosphate buffer, phosphate-citrate buffer, and succinate buffer.

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- 21. The formulation of claim 16, wherein said botulinum toxin is of a botulinum toxin type selected from the group consisting of Types A, B, C₁, C₂, D, E, F and G.
- 22. The formulation of claim 21, wherein said botulinum toxin is botulinum toxin Type B present at a concentration of about 100-20,000 U/ml.
- 23. The formulation of claim 22, wherein said botulinum toxin Type B is present in a high molecular weight complex of about 700 kD.
- 24. The formulation of claim 22, wherein said botulinum toxin Type B is present at a concentration in the range of about 1000-5000 U/ml.
- 25. The formulation of claim 21, wherein said botulinum toxin is botulinum toxin Type A, present at a concentration in the range of about 20-2000 U/ml.
- 26. The formulation of claim 25, wherein said botulinum toxin is botulinum toxin Type A, present at a concentration in the range of about 100-1000 U/ml.
 - 27. The formulation of claim 16, which further includes an excipient protein.
- 28. The formulation of claim 25, wherein said excipient protein is selected from the group consisting of serum albumin, human serum albumin, and gelatin.